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=> s fagales

L1143 FAGALES

=> s recombinant

L2 622962 RECOMBINANT

=> s 12 and fagales

46 L2 AND FAGALES T.3

=> s 13 and substitution

0 L3 AND SUBSTITUTION

=> s 13 and amino acid substitution

L5 0 L3 AND AMINO ACID SUBSTITUTION

=> s recombinant allergen

L6 1058 RECOMBINANT ALLERGEN.

=> s 16 and V bet1

L7 0 L6 AND V BET1

=> s 16 and Bet v1

6 L6 AND BET V1

=> s 18 and amino acid substitution

0 L8 AND AMINO ACID SUBSTITUTION

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ANSWER 1 OF 6 BIOSIS COPYRIGHT 2001 BIOSIS

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DN PREV199900330765

How far can we simplify in vitro diagnostics for Fagales tree pollen allergy? A study with three whole pollen extracts and purified natural

and

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recombinant allergens.
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AU van Ree, R. (1); van Leeuwen, W. A.; Akkerdaas, J. H.; Aalberse, R. C. CS (1) Department of Allergy, C.L.B., Plesmanlaan 125, NL-1066 CX, Amsterdam Netherlands

SO Clinical and Experimental Allergy, (June, 1999) Vol. 29, No. 6, pp. 848-855.

ISSN: 0954-7894.

DT Article

LA English

SL English

AB Background Current diagnostic tests for Fagales tree pollen allergy are often composed of mixtures of pollen of birch, alder and hazel. Their complex composition hampers accurate standardization. Objective The aim of

this study was to investigate whether mixtures of tree pollen extracts can

be replaced by a single pollen species, and whether a single pollen species can be replaced by a limited number of purified natural or recombinant major allergens. Methods Sera (n = 1725) were selected on ground of a general suspicion for inhalant allergy, and tested in a RAST for birch, alder and hazel pollen. Sera with > 0.5 RU/mL for any of the three species were tested in a RAST for natural Bet v 1 and Bet v 2 as well as for recombinant versions of both allergens. Results Specific IgE antibodies (> 0.3 RU/mL) against birch, alder and hazel were found in

298 and 292 sera, respectively. All sera with a positive RAST for alder and/or hazel and a negative RAST for birch were low-responder sera on alder and hazel, only five sera having a RAST value > 1.0 (all < 2.0).

For

242,

all sera with a RAST  $> 0.5 \ \text{RU/mL}$  (n = 250), the mean of individual ratio's

alder/birch and hazel/birch was 1.02 and 0.54, respectively. Of 223 of these sera, 63.2% had specific IgE against natural Bet v 1 and 63.7% against natural Bet v 2. When responses to both allergens were combined 93.7% were positive. The mean ratios Bet v 1 +  $2/\exp(1.00)$  and 2.11 in case of birch, alder and hazel, respectively. For 211 sera

the six

same analysis was performed with recombinant Bet v 1 and Bet v 2. Only

sera with Bet v 1-specific IgE (all < 0.5 RU/mL) were negative (< 0.3 RU/mL) on recombinant Bet v 1. For Bet v 2, 77/132 sera with specific IgE to the natural allergen did not react to the recombinant version. Twelve false-negatives had RAST values > 1.0 RU/mL. The mean of the individual recombinant/natural ratios was 0.98 for Bet v 1 and 0.38 for Bet v 2 (P < 0.001). The mean ratio rBet v 1 + 2/birch was 0.75 with 17.5% false-negatives on the combination of  $\bf recombinant$ 

**allergens**. Conclusion Reliable in vitro diagnosis is possible with a single tree pollen extract (birch or alder). The same is true for purified natural Bet v 1 and Bet v 2. A combination of recombinant molecules is slightly less efficient.

CC Allergy \*35500

Pathology, General and Miscellaneous - Diagnostic \*12504 Plant Physiology, Biochemistry and Biophysics - Reproduction \*51512 Respiratory System - General; Methods \*16001

BC Hominidae 86215

IT Major Concepts

Allergy (Clinical Immunology, Human Medicine, Medical Sciences); Pulmonary Medicine (Human Medicine, Medical Sciences)

IT Diseases

Feagles tree pollen allergy: immune system disease, respiratory system disease  $% \frac{1}{2}\left( \frac{1}{2}\right) =\frac{1}{2}\left( \frac{1}{2}\right) +\frac{1}{2}\left( \frac{1}{2}\right) +\frac{1}{2}\left($ 

IT Chemicals & Biochemicals

Bet v1: allergen; Bet v2: allergen

```
Miscellaneous Descriptors
        diagnosis
ORGN Super Taxa
        Hominidae: Primates, Mammalia, Vertebrata, Chordata, Animalia
ORGN Organism Name
        human (Hominidae)
ORGN Organism Superterms
        Animals; Chordates; Humans; Mammals; Primates; Vertebrates
^{\text{L8}}
     ANSWER 2 OF 6 BIOSIS COPYRIGHT 2001 BIOSIS
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TI
     Molecular characterization of Dau c 1, the Bet v 1 homologous protein
from
     carrot and its cross-reactivity with Bet v 1 and Api g 1.
     Hoffmann-Sommergruber, K. (1); O'Riordain, G.; Ahorn, H.; Ebner, C.; Da
ΑU
     Camara Machado, M. Laimer; Puehringer, H.; Scheiner, O.; Breiteneder, H.
     (1) Department of General and Experimental Pathology, AKH-EBO 3Q,
CS
     Waehringer Guertel 18-20, A-1090, Vienna Austria
SO
     Clinical and Experimental Allergy, (June, 1999) Vol. 29, No. 6, pp.
     840-847.
     ISSN: 0954-7894.
DT
     Article
LA
     English
SL
     English
AΒ
     Background Up to 70% of patients with birch pollen allergy exhibit the
     so-called oral allergy syndrome, an IgE-mediated food allergy. The most
     frequent and therefore best characterized pollen-fruit syndrome is apple
     allergy in patients suffering from tree pollen-induced pollinosis. The
     occurrence of adverse reactions to proteins present in vegetables such as
     celery and carrots in patients suffering from pollen allergy has also
been
     reported. cDNAs for Bet v 1 homologous proteins have been cloned from
     celery, apple and cherry. Objective The aim of the study was to identify
     Bet v 1 homologues from carrot (Daucus carota), to isolate the respective
     cDNA, to compare the IgE-binding capacity of the natural protein to the
     recombinant allergen and determine the cross-reactivity
     to Api g 1 and Bet v 1. Methods Molecular characterization of the carrot
     allergen was performed using IgE-immunoblotting, cross-inhibition assays,
     N-terminal sequencing, PCR-based cDNA cloning and expression of the
     recombinant protein in Escherichia coli. Results A 16-kDa protein from
     carrot was identified as a major IgE-binding component and designated Dau
     c 1. Sequencing corresponding cDNAs revealed three extremely similar
     sequences (Dau c 1.1, 1.2 and 1.3) with an open reading frame of 462 bp
     coding for 154 amino acid residues. Conclusions Purified recombinant Dau
     1.2 was tested in immunoblots displaying IgE-binding capacity comparable
     to its natural counterpart. Cross-inhibition assays verified the
existence
     of common B-cell epitopes present on Dau c 1, Api g 1 as well as on Bet v
CC
     Allergy *35500
     Biochemical Studies - General *10060
     Plant Physiology, Biochemistry and Biophysics - Reproduction *51512
     Respiratory System - General; Methods *16001
BC
     Hominidae
                 86215
ΙT
    Major Concepts
        Allergy (Clinical Immunology, Human Medicine, Medical Sciences);
        Respiratory System (Respiration)
IΤ
     Diseases
       birch pollen allergy: immune system disease, respiratory system
```

disease IT Ch

Chemicals & Biochemicals

birch pollen: allergen; cDNA [complementary DNA]; Dau cl protein:
allergen, carrot Bet V1 protein

ORGN Super Taxa

Hominidae: Primates, Mammalia, Vertebrata, Chordata, Animalia

ORGN Organism Name

human (Hominidae): patient

ORGN Organism Superterms

Animals; Chordates; Humans; Mammals; Primates; Vertebrates

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- AN 1997:243013 BIOSIS
- DN PREV199799542216
- TI Cross-reacting allergens in tree pollen and pollen-related food allergy: Implications for diagnosis of specific IgE.
- AU Scheiner, O. (1); Aberer, W.; Ebner, C.; Ferreira, F.; Hoffmann-Sommergruber, K.; Hsieh, L. S.; Kraft, D.; Sowka, S.; Vanek-Krebitz, M.; Breiteneder, H.
- CS (1) Dep. Gen. Experimental Pathol., AKH-EBO 3Q, Waehringer Guertel 18-20, A-Vienna Austria
- SO International Archives of Allergy and Immunology, (1997) Vol. 113, No. 1-3, pp. 105-108.
  ISSN: 1018-2438.
- DT Article
- LA English
- AB Background: A number of recombinant allergens are by